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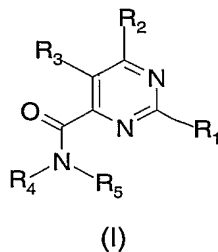
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(54) Title: PYRIMIDINE COMPOUNDS AS PURINE RECEPTOR ANTAGONIST



(57) Abstract: Compounds of formula (I); wherein R<sub>1</sub> is H or NHZ; R<sub>2</sub> is optionally substituted aryl or heteroaryl attached via a carbon atom; R<sub>3</sub> is H; optionally substituted C<sub>1</sub>-C<sub>6</sub>alkyl, C<sub>2</sub>-C<sub>6</sub>alkenyl, C<sub>2</sub>-C<sub>6</sub>alkynyl, or C<sub>3</sub> C<sub>7</sub> cycloalkyl, halogen; OH or OR, or R<sub>4</sub> is H, optionally substituted C<sub>1</sub>-C<sub>6</sub>alkyl, C<sub>3</sub> C<sub>6</sub>alkenyl, C<sub>3</sub>-C<sub>6</sub>alkynyl, C<sub>3</sub>-C<sub>7</sub> cycloalkyl, aryl or heteroaryl, R<sub>5</sub> is H or optionally substituted C<sub>1</sub>-C<sub>6</sub>alkyl, C<sub>3</sub>-C<sub>6</sub>alkenyl, C<sub>3</sub>-C<sub>6</sub>alkynyl, or C<sub>3</sub>-C<sub>7</sub> cycloalkyl; or R<sub>4</sub> and R<sub>5</sub> together form a 5 or 6-membered heterocyclic ring; and R<sub>10</sub> is optionally substituted C<sub>1</sub>-C<sub>6</sub>alkyl; are purine receptor, particularly adenosine receptor antagonists, useful for treatment of, inter alia, movement disorders such as Parkinsons disease.

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